	UNITED KINGDOM	Animal health/Official certificate to the EU				
	I.1 Consignor/Exporter		I.2 Certificate ref	ference	I.2a	
	Name					
	Address		I.3 Central Comp	npetent Authority		
			DEPARTMENT F FOOD & RURAL	FOR ENVIRONMENT, L AFFAIRS		
′			I.4 Local Compet	tent Authority	1 /	
	Country	ISO country code	ANIMAL AND P	LANT HEALTH AGENCY		
	1.5 Consignee/Importer	I.6 Operator resp	onsible for the consignment	t		
	Name		Name			
4	Address	Address				
Part I: Description of consignment	· 6					
0 u 0	Country	ISO country code	Country	IS	SO country code	
scriptio	I.7 Country of origin	ISO country code	I.9 Country of de	stination	ISO country code	
t I: De	I.8 Region of origin Code		I.10 Region of des	of destination Code		
Par	I.11 Place of dispatch	Registration/Approval No	I.12 Place of desti	ination	Registration/Approval No	
ĺ	Name		Name			
	Address		Address			
	Country	ISO country code	Country	I:	SO country code	
	I.13 Place of loading	ibo country is a	I.14 Date and tim		50 county 1111	
			U			
	I.15 Means of transport	I.16 Entry Border Control Post				
l	□ Aircraft □ V	Vessel	I.17 Accompanyi	ng documents		
		ļ	Туре	Co	.Ao	
	□ Railway □ R	Road vehicle				
	□ Ranway □ F	toad venicle	Country	ISO	O country code	
	T1 - 20 - 2		Commercial docur	ment reference		
	Identification					
	I.18 Transport conditions	□ Ambient	□ Chilled	□ Froze	en	
	I.19 Container number/Seal number					
	Container No	Seal No				
	I.20 Certified as or for					
	□ Products for human consumpti					
	I.21		I.22			
	Third country	ISO country code	I.23			
Ī	I.24 Total number of packages	I.25		I.26 Total net weight/gross	s weight (kg)	

	NA.				II.a Certificate refer	
	OM of consignment					
CN Code	Species	Nature of	Number of	Type of packaging	Batch No	Net Weigh
		commodity	packages	71 - 1		
Treatment Type	Date of collection/production	Manufacturing plant	Cold Store			Final consumer
* *						
CN Code	Species	Nature of commodity	Number of packages	Type of packaging	Batch No	Net Weigh
Treatment Type	Date of collection/production	Manufacturing plant	Cold Store			Final consumer □
1		<u> </u>				
CN Code	Species	Nature of commodity	Number of packages	Type of packaging	Batch No	Net Weigh
Treatment Type	Date of collection/production	Manufacturing plant	Cold Store	0		Final consumer □
CN Code	Species	Nature of commodity	Number of packages	Type of packaging	Batch No	Net Weigh
Treatment Type	Date of collection/production	Manufacturing plant	Cold Store		C/1	Final consumer
<u> </u>						
CN Code	Species	Nature of commodity	Number of packages	Type of packaging	Batch No	Net Weigh
Treatment Type	Date of collection/production	Manufacturing plant	Cold Store			Final consumer

II.a Certificate reference

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II. Health information

II.1. Public health attestation [Delete when the Union is not the final destination of the dairy products]

the undersigned, declare that I am aware of the relevant requirements of Regulation (EC) No 178/2002 of the European Parliament and of the Council, Regulation (EC) No 852/2004 of the European Parliament and of the Council, Regulation (EC) No 853/2004 of the European Parliament and of the Council and Regulation (EU) 2017/625 of the European Parliament and of the Council and Commission Implementing Regulation (EU) 2019/627 and hereby certify that the dairy product described in Part I was produced in accordance with these requirements, and in particular that:

- it was produced from raw milk:
 - which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Articles 49 and 50 of Implementing Regulation (EU) 2019/627;
 - which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Section IX, Chapter I, of Annex III to Regulation (EC) No 853/2004;
 - (iii) which meets the plate and somatic cell count criteria laid down in Section IX, Chapter I, of Annex III to Regulation (EC) No 853/2004:
 - which complies with the guarantees covering live animals and products thereof provided by the control plan submitted in accordance (iv) with Article 6(2) of Commission Delegated Regulation (EU) 2022/2292 and the third country or region thereof of its origin is listed in Annex - I to Commission Implementing Regulation (EU) 2021/405 with an entry 'X' for milk;
 - which, pursuant to testing for antibiotic residues carried out by the food business operator in accordance with the requirements (v) Section IX, Chapter I, Part III, point 4, of Annex III to Regulation (EC) No 853/2004, complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Commission Regulation (EU) No 37/2010;
 - has not been obtained from animals showing a positive reaction to the test for tuberculosis or brucellosis; (vi)
- it comes from (an) establishment(s) applying general hygiene requirements and implementing a programme based on the hazard analysis and (b) critical control points (HACCP) principles in accordance with Article 5 of Regulation (EC) No 852/2004, regularly audited by the competent authorities, and is listed as a Union approved establishment;
- (c) it has been processed, stored, wrapped, packaged and transported in accordance with the relevant hygiene conditions laid down in Annex II to Regulation (EC) No 852/2004 and Section IX, Chapter II, of Annex III to Regulation (EC) No 853/2004;
- it meets the relevant criteria laid down in Section IX, Chapter II, of Annex III to Regulation (EC) No 853/2004 and the relevant microbiological criteria laid down in Commission Regulation (EC) No 2073/2005;
- it has undergone or been produced from raw milk which has been submitted to a treatment involving a single heat treatment with a heating (e) effect at least equivalent to that achieved by a pasteurisation process of at least 72°C for 15 seconds and, where applicable, sufficient to ensure a negative reaction to an alkaline phosphatase test immediately after the heat treatment.
- Attestation as regards Commission Delegated Regulation (EU) 2023/905 [Delete when the Union is not the final (1)(4)[II.1.a. destination of the dairy products]

I, the undersigned, declare that I am aware of the relevant requirements of Regulation (EU) 2019/6 of the European Parliament and of the Council and Commission Delegated Regulation (EU) 2023/905 and hereby certify that the dairy product described in Part I was produced in accordance with these requirements, and in particular that, the animals from which the raw milk has been obtained have not been administered antimicrobial medicinal products for growth promotion or yield increase or antimicrobial medicinal products containing an antimicrobial that is included in the list of antimicrobials reserved for the treatment of certain infections in humans laid down in Commission Implementing Regulation (EU) 2022/1255 as set out in Article 3 of Delegated Regulation (EU) 2023/905 and originate from a third country or region thereof listed in accordance with Article 5(2) of Delegated Regulation (EU) 2023/905.]

Animal health attestation [Delete when the dairy products are derived from solipeds, leporidae or wild land mammals other than ungulates] II.2.

The **dairy products** described in Part I:

- for the entry into the Union of milk and listed in Part 1 of Annex XVII to Commission Implementing Regulation (EU) 2021/404, and in which foot and mouth disease and infection with rinderpest virus have not been reported for the last 12 months prior to the date of milking, and vaccination against these diseases has not been carried out during that period;
- II.2.2. have been processed from:

(1) either [II.2.2.1. raw milk originating from:

 $^{(1)}$ either [the zone referred to in point II.2.1. and obtained from animals of the species [Bos taurus,] (1) [Ovis aries,] (1) [Capra hircus,] (1) [Bubalus bubalis,] (1) [Camelus dromedarius] (1) that:

- (1) either have remained in the zone referred to under point II.2.1. since birth, or for the last three months prior to the date of milking;]
- (1) and/or [(a) were introduced in the zone referred to under point II.2.1. from:

(1) either

[another third country or territory, or zone thereof which is listed for entry into the Union of milk, colostrum or colostrum-based products and the animals remained there for the last three months prior to the date of milking;]]

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II.a Certificate reference

(1) and/or [a Member State;]]

(b) have been kept in **establishments**:

- registered by and under the control of the competent authority of the third country or territory and have a system in place to maintain and to keep records in accordance with Article 8 of Commission Delegated Regulation (EU) 2020/692;
- (ii) which receive regular animal health visits from a veterinarian for the purpose of the detection of, and information on, signs indicative of the occurrence of diseases, including the listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 relevant for the species and emerging diseases;
- (iii) which were not subject to national restriction measures for animal health reasons, including the listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 relevant for the species and emerging diseases, at the date of milking.]

(1) and/or [a Member State.]

(1) and/or [II.2.2.1 dairy products

(a) produced in

(1) either [the zone referred to in point II.2.1.;]]

Union as such upon arrival in the zone referred to under point II.2.1.]]

(1) and/or [a Member State.]]

(b) obtained **from raw milk** originating from:

(1) either [the zone referred to in point II.2.1. and obtained from animals of the species [Bos taurus,] (1) [Ovis aries,] (1) [Capra hircus,] (1) [Bubalus bubalis,] (1) [Camelus dromedarius] (1) that:

(1) either [(i) have remained in the zone referred to under point II.2.1. since birth, or for the last three months prior to the date of milking;]

 $^{(1)}$ and/or [(i) were introduced in the zone referred to under point II.2.1. from:

(1) either [another third country or territory, or zone thereof which is listed for entry into the Union of milk, colostrum or colostrum-based products and the animals remained there for the last three months prior to the date of milking;]]

(1) and/or [a Member State;]]

(ii) have been kept in **establishments**:

- registered by and under the control of the competent authority of the third country or territory and have a system in place to maintain and to keep records in accordance with Article 8 of Delegated Regulation (EU) 2020/692;
- (ii) which receive regular animal health visits from a veterinarian for the purpose of the detection of, and information on, signs indicative of the occurrence of diseases, including the listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 relevant for the species and emerging diseases;
- (iii) which were not subject to national restriction measures for animal health reasons, including the listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 relevant for the species and emerging diseases, at the date of milking.]]

(1) and/or [a Member State.]]

Notes

In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, references to the Union in this animal health/official certificate include the United Kingdom in respect of Northern Ireland.

This animal health/official certificate is intended for the entry into the Union of dairy products (as defined in point 7.2 of Annex I to Regulation (EC) No 853/2004) entering from zones listed in Annex XVII to Implementing Regulation (EU) 2021/404 for entry into the Union of milk and therefore not

II.a Certificate reference

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required to undergo a specific risk-mitigating treatment against foot and mouth disease but are required to undergo a pasteurisation treatment because they were produced from raw milk obtained in the establishments which are not officially free of tuberculosis or free or officially free of brucellosis, including when the Union is not the final destination of such dairy product.

This animal health/official certificate shall be completed in accordance with the notes for the completion of certificates provided for in Chapter 4 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

Box reference I.8.: Provide the code of the zone as appearing in column 2 of the table in Part 1 of Annex XVII to Implementing Regulation

(EU) 2021/404.

Box reference I.11.: Name, address and approval number of the establishment of dispatch.

Box reference I.1 Registration number (railway wagons or container and road vehicles), flight number (aircraft) or name (vessel) must be provided. In the case of transport in containers their registration number and where there is a serial number of the seal it

must be indicated in Box I.19. In the case of unloading and reloading, the consignor must inform the border control post

of entry into the Union.

Box reference I.19 For the containers or boxes, the container number and the seal number (if applicable) shall be included.

Box reference I.27 .: Description of consignment:

> code": Indicate the appropriate Harmonised System (HS) code(s) of the World Customs Organisation under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 15.17; 17.02; 21.05; 21.06; 28.35; 35.01; 35.02 or 35.04.

> "Manufacturing plant": Introduce the approval number of the treatment and/or processing establishment(s) approved for

the entry into the Union

Part II:

Delete if not applicable.

(2) Code of the zone in accordance with column 2 of the table in Part 1 of Annex XVII to Implementing Regulation (EU) 2021/404.

(3) to be signed by:

an official veterinarian when Part II.2. Animal health attestation is not deleted,

a certifying officer or an official veterinarian when Part II.2. Animal health attestation is deleted.

(4) Applicable to consignments entering the Union as from 3 September 2026.

[Official veterinarian] (1) (3)/[Certifying officer] (1) (3)

Name (in capital letters)

Date Qualification a

Stamp Signature